Supplemental Table 6. Adverse Events

	Pre-Randomization (N=10)	During AID Use Period (N=35)	During SAP Use Period (N=35)
Any Reportable Adverse Event[a]			
Number of Events	1	6	1
Participants with ≥ 1 Event	1 (10%)	5 (14%)	1 (3%)
Specific Events – # *Participants [# Events]*			
Hyperglycemia	0 [0]	3 [3]	0 [0]
Hypoglycemia	0 [0]	0 [0]	1 [1]
Ketosis	0 [0]	1 [1]	0 [0]
Medical Device Site Bleeding	0 [0]	1 [1]	0 [0]
COVID-19	1 [1]	1 [1]	0 [0]
Days with Ketone Values ≥1.5 mmol/L		1	0

[a] Reportable adverse events include:

- An SAE
- An adverse device effect (ADE)
- An adverse event occurring in association with study procedure
- An adverse event which leads to the discontinuation of a study device for two or more hours
- Hypoglycemia meeting the definition of severe hypoglycemia
- Diabetic ketoacidosis